

# Genome Sequence of the Persistent *Salmonella enterica* subsp. *enterica* Serotype Senftenberg Strain SS209

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*Salmonella enterica* subsp. *enterica* serotype Senftenberg is an emerging serotype in poultry production which has been found to persist in animals and the farm environment. We report the genome sequence and annotation of the SS209 strain of *S. Senftenberg*, isolated from a hatchery, which was identified as persistent in broiler chickens.

**S**almonellosis are a worldwide health problem and are usually associated with poultry products. Over the last decade, the emergence of some serotypes in poultry production has been observed (1, 14). *Salmonella enterica* serotype Senftenberg has always been associated with the hatchery environment, but recently it has become more frequent in poultry farms (7, 10). Furthermore, some strains were responsible for several human infections throughout the world (8, 11). In order to understand the emergence of the Senftenberg serotype in poultry, we previously identified a group of *S. Senftenberg* strains showing high intestinal colonization levels and able to persist over several weeks in broiler chickens, contrary to the case with other strains (2). The SS209 strain presented in this study is one of the persistent strains. It originated with 1-day-old chicks and belongs to the multilocus sequence type (MLST) ST14, the most common sequence type found in serotype Senftenberg worldwide (13).

Here we report the genome sequence of *S. Senftenberg* SS209, obtained using a combination of 454 pyrosequencing and Illumina genome analyzer IIx paired-end reads (performed by GATC Biotech, Konstanz, Germany). Sequencing yielded 239,453 reads for 454 sequencing (17.2-fold coverage) and 1,733,213 paired-end reads for Illumina (24.8-fold coverage), which were assembled *de novo* using the Newbler 2.3 assembly software program (Roche). The minimum contig size was set to 500 nucleotides (nt), which generated 109 contigs for the bacterial chromosome. No sequences corresponding to the large plasmid present in some *Salmonella* serotypes were detected. Genome annotation was performed using the AGMIAL annotation platform (3) and the *S. Typhimurium* strain LT2 as a reference (9).

The chromosome of *S. Senftenberg* SS209 has an overall G+C content of 51.73% and a predicted genome size of about 5.02 Mb, which is similar to the estimated genome length of the other sequenced *S. Senftenberg* strain (6). It is composed of 4,838 coding sequences (CDS). A pseudogenome of SS209 has been constructed by ordering contigs using the Mauve Contig Mover software program (12) and the LT2 strain genome as a reference. Genomic alignment between SS209 and other *Salmonella* serotypes was then performed using the Mauve genome aligner (5) and postprocessed for integration into the MOSAIC resource (4). A first analysis of whole-genome comparisons showed that *S. Senftenberg* SS209 is quite distant from the serotypes commonly found in poultry, namely, *Salmonella enterica* serotypes Typhimurium, Enteritidis, and Gallinarum. The core genome (i.e., back-

bone) common to *S. Senftenberg* SS209 and any of the three other serotypes ranges between 4.20 Mb (with *S. Gallinarum* 287/91), 4.24 Mb (with *S. Enteritidis* P125109), and 4.33 Mb (with *S. Typhimurium* LT2) and represents, respectively, 80.7% (*S. Gallinarum*), 81.4% (*S. Enteritidis*), and 83.1% (*S. Typhimurium*) of the SS209 genome. The aligned pairs of backbones share 98.8% of identity on average compared to any of the *S. Gallinarum*, *S. Typhimurium*, and *S. Enteritidis* genomes. Further genomic studies should allow a better understanding of *S. Senftenberg* evolution, especially the relationships between this emergent serotype and the other pathogenic *Salmonella* serotypes.

**Nucleotide sequence accession number.** The *S. Senftenberg* strain SS209 whole-genome sequence assembly and its annotation have been deposited in EMBL under the project accession number [CAG000000000](#).

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## REFERENCES

1. Anonymous. 2010. Analysis of the baseline survey on the prevalence of *Campylobacter* in broiler batches and of *Campylobacter* and *Salmonella* on broiler carcasses in the EU, 2008. Part A: *Campylobacter* and *Salmonella* prevalence estimates. EFSA J. 8:1503.
2. Boumart Z, et al. Heterogeneity of persistence of *Salmonella enterica* serotype Senftenberg strains could explain the emergence of this serotype in poultry flocks. PLoS One, in press.
3. Bryson K, et al. 2006. AGMIAL: implementing an annotation strategy for prokaryote genomes as a distributed system. Nucleic Acids Res. 34:3533–3545.
4. Chiappello H, et al. 2008. MOSAIC: an online database dedicated to the comparative genomics of bacterial strains at the intra-species level. BMC Bioinformatics 9:498.
5. Darling ACE, Mau B, Blattner FR, Perna NT. 2004. Mauve: multiple

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- alignment of conserved genomic sequence with rearrangements. *Genome Res.* 14:1394–1403.
6. den Bakker HC, et al. 2011. Genome sequencing reveals diversification of virulence factor content and possible host adaptation in distinct subpopulations of *Salmonella enterica*. *BMC Genomics* 12:425.
  7. Hamada S, Hashimoto H, Tasaka T, Tsuchiya Y. 1958. Studies on chick salmonellosis. II. *Salmonella* Senftenberg infection in chicks. *Jpn. J. Vet. Res.* 6:181–195.
  8. Hu Q, et al. 2008. *Salmonella enterica* serovar Senftenberg human clinical isolates lacking SPI-1. *J. Clin. Microbiol.* 46:1330–1336.
  9. McClelland M, et al. 2001. Complete genome sequence of *Salmonella enterica* serovar Typhimurium LT2. *Nature* 413:852–856.
  10. Pedersen TB, Olsen JE, Bisgaard M. 2008. Persistence of *Salmonella* Senftenberg in poultry production environments and investigation of its resistance to desiccation. *Avian Pathol.* 37:421–427.
  11. Pezzoli L, et al. 2007. International outbreak of *Salmonella* Senftenberg in 2007. *Euro Surveill.* 12:pii=3218.
  12. Rissman AI, et al. 2009. Reordering contigs of draft genomes using the Mauve Aligner. *Bioinformatics* 25:2071–2073.
  13. Stepan RM, Sherwood JS, Petermann SR, Logue CM. 2011. Molecular and comparative analysis of *Salmonella enterica* Senftenberg from humans and animals using PFGE, MLST and NARMS. *BMC Microbiol.* 11:153.
  14. Voetsch AC, et al. 2004. FoodNet estimate of the burden of illness caused by nontyphoidal *Salmonella* infections in the United States. *Clin. Infect. Dis.* 38(Suppl. 3):S127–S134.